

PCTWORLD INTELLECTUAL PROPERTY ORGANIZATION
International Bureau

INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification 5 : C07C 2/12	A2	(11) International Publication Number: WO 93/16020 (43) International Publication Date: 19 August 1993 (19.08.93)
(21) International Application Number: PCT/EP93/00177 (22) International Filing Date: 27 January 1993 (27.01.93) (30) Priority data: 92300797.5 30 January 1992 (30.01.92) EP (34) Countries for which the regional or international application was filed: DE et al. (71) Applicant (for all designated States except US): EXXON CHEMICAL PATENTS INC. [US/US]; 1900 East Linden Avenue, Florham Park, NJ 07036 (US). (72) Inventors; and (75) Inventors/Applicants (for US only) : MATHYS, Georges, Marie, Karel [BE/BE]; Eikenboomlaan 66, B-3360 Bierbeek (BE). MARTENS, Luc, Roger, Marc [BE/BE]; Prisenbeemdlaan 3, B-1860 Meise (BE). BAES, Marleen, Augusta [BE/BE]; Don Boscolaan 11, B-3050 Oud-Heverlee (BE). VERDUIJN, Johannes, Petrus [NL/NL]; Westersingel 34, NL-3202 XL Spijkinesse (NL). HUYBRECHTS, Diane, Renata, Cornelia [BE/BE]; Burgemeester Van Loonstraat 4, B-2300 Oud-Turnhout (BE).		(74) Agents: BAWDEN, Peter, Charles et al.; Exxon Chemical Limited, Exxon Chemical Technology Centre, P.O. Box 1, Abingdon, Oxfordshire OX13 6BB (GB). (81) Designated States: CA, JP, US, European patent (AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE). Published <i>Without international search report and to be republished upon receipt of that report.</i>
(54) Title: ALKENE OLIGOMERIZATION (57) Abstract In a process for the oligomerization of C ₂ -C ₁₂ -alkenes, an alkene-containing feedstock having a water content of from 0.05 to 0.25 molar %, based on the hydrocarbon content of the feedstock, is passed over a zeolite catalyst. Improved alkene conversion and catalyst life are obtained compared to processes in which non-hydrated feedstocks are used.		

FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AT	Austria	FR	France	MR	Mauritania
AU	Australia	GA	Gabon	MW	Malawi
BB	Barbados	GB	United Kingdom	NL	Netherlands
BE	Belgium	GN	Guinea	NO	Norway
BF	Burkina Faso	GR	Greece	NZ	New Zealand
BG	Bulgaria	HU	Hungary	PL	Poland
BJ	Benin	IE	Ireland	PT	Portugal
BR	Brazil	IT	Italy	RO	Romania
CA	Canada	JP	Japan	RU	Russian Federation
CF	Central African Republic	KP	Democratic People's Republic of Korea	SD	Sudan
CC	Congo	KR	Republic of Korea	SE	Sweden
CH	Switzerland	KZ	Kazakhstan	SK	Slovak Republic
CI	Côte d'Ivoire	LI	Liechtenstein	SN	Senegal
CM	Cameroon	LK	Sri Lanka	SU	Soviet Union
CS	Czechoslovakia	LU	Luxembourg	TD	Chad
CZ	Czech Republic	MC	Monaco	TG	Togo
DE	Germany	MG	Madagascar	UA	Ukraine
DK	Denmark	ML	Mali	US	United States of America
ES	Spain	MN	Mongolia	VN	Viet Nam
FI	Finland				

ALKENE OLIGOMERISATION

This invention relates to the oligomerisation of alkenes and to catalysts used in alkene oligomerisation processes in particular for the production of C₅ to C₁₅ olefins which may be subsequently subject to hydroformylation for the production of aldehydes and alcohols themselves useful in the production of plasticiser esters, mercaptans and surfactants.

In one form of olefin oligomerisation a solid phosphoric acid catalyst is used as catalyst. A significant amount of cracked products is produced, the phosphoric acid catalyst is not regenerable and its disposal in an environmentally acceptable manner is difficult and expensive.

Zeolite catalysts of many types have been proposed for the conversion of lower molecular weight olefins (that is, olefins having up to twelve carbon atoms) into olefinic hydrocarbons of higher molecular weight. The higher molecular weight olefins may be used in a variety of chemical processes, for example, in the manufacture of fuel oil, synthetic lubricants, alcohols, plasticisers and mercaptans.

International Patent Application WO 89/07586 discloses a process for conversion of a fuel gas containing ethene and propene and a catalytic reformat containing C₆ to C₈ aromatic hydrocarbons into hydrocarbon products of higher molecular weight. The fuel gas and reformat feedstocks are contacted at elevated temperature and pressure with a zeolite catalyst, which may be of the ZSM-5 type. If ZSM-5 pentasil type crystals are used, the catalyst crystals preferably have a size of from 0.01 to over 2 microns. WO 89/07586 indicates that the zeolites ZSM-22 and ZSM-23 may be used.

US Patent 4868146 discloses a catalyst having a crystalline metallosilicate inner core and an inactive porous shell, which may be used, for example, to catalyse the oligomerisation of

lower olefins. Highly siliceous zeolites, including, for example, ZSM-22, may be used. US Patent 4788374 similarly discloses an olefin oligomerisation catalyst having a crystalline metallosilicate core and a porous shell.

5 European Specification 377306 discloses a process in which branched internal olefins are co-oligomerised with alpha-olefins to give a synthetic lube-range product. In a first step, propene is converted by contact with a zeolite catalyst
10 of the ZSM-5 type to a mixture of oligomers. In a number of the Examples a ZSM-23 catalyst that has been prepared in accordance with US Patent 4160788 is used.

15 European Specifications 276095, 323032 and 318132 disclose processes for oligomerising light olefins using zeolite catalysts.

US Patent 4481177 discloses the use of ZSM-22 in the organic nitrogen-containing and alkali-metal containing form. That
20 form is apparently obtained by cation exchange of the ZSM-22 with an alkyl- or aryl-ammonium ion. US 4481177 also suggests that ZSM-22 may be used to catalyse various chemical reactions, including olefin oligomerisation. There is no Example, however, of any olefin oligomerisation.

25 European Specification 0299671 discloses the production of high viscosity index lubricating oils by the oligomerisation of lower olefins over ZSM-5 zeolite catalysts by cofeeding small amounts of water with the hydrocarbon stream. The
30 lubricating oils have a boiling point above 343°C and it is speculated that the presence of the water enhances the intracrystalline acid site polymerisation in preference to surface active site polymerisation leading to the formation of more linear lubricant range hydrocarbons and thus an
35 enhancement in viscosity index.

It has now been found that, unexpectedly, by using in an alkene oligomerisation process an alkene-containing feedstock with a water content of from 0.05 to 0.25 molar %, and preferably of at least 0.06 molar %, based on the hydrocarbon content of the feedstock, the yields of the desired higher molecular weight alkenes can be increased, and the catalyst becomes deactivated more slowly.

The invention provides a process for improving the yield and prolonging catalyst life in oligomerising at least one alkene having from 2 to 12 carbon atoms, (hereinafter referred to for simplicity as "C₂-C₁₂-alkenes") comprising contacting a C₂-C₁₂-alkene-containing feedstock having a water content of 0.05 to 0.25 molar %, based on the hydrocarbon content of the feedstock, with a zeolite catalyst.

The invention is particularly concerned with the production of C₅ to C₂₀ olefins of initial boiling point 30 to 310°C, preferably 30 to 300°C, more preferably 30 to 250°C, from propylene and/or butene feedstocks. In particular the production of the following olefins.

Products	Distillation Range, °C	
	ASTM D1078	
	Initial Boiling Point	Dry Point
Pentenes	30	
Hexenes	63	
Heptenes	88	97
Octenes	114	126
Nonenes	135	143
Decenes	155	160
Undecenes	167	178
Dodecenes	185	194
Tetramer K	181	200
Tetramer V	186	198
Tetramer D	186	224
Tetramer P	189	225
Tridecenes	204	213

In addition to enabling the yields of higher molecular weight alkenes to be increased, the process of the invention enables the oligomerisation reaction to be carried out at relatively low temperatures. Preferably, the contacting of the feedstock with the catalyst is carried out at a temperature not exceeding 260°C. Advantageously, the said contacting is carried out at a temperature of from 180 to 255°C.

In the case of an alkene-containing feedstock having a water content of less than 0.05 molar %, the water content may be increased by any suitable means. For example, the feedstock can be passed through a thermostatted water saturator. Since the amount of water required to saturate the alkene feedstock will depend upon the temperature of the feedstock, control of the water content can then be effected by appropriate control of the temperature of the feedstock. The water content of the feedstock is preferably at least 0.06 molar %, based on the hydrocarbon content of the feedstock.

The zeolite catalyst may be any catalyst that is active in alkene oligomerisation reactions. For example, there may be used a catalyst selected from the group consisting of zeolites of the TON structure type (for example, H-ZSM-22, H-ISI-1, H-Theta-1, H-Nu-10, KZ-2), or zeolites of the MTT structure type (for example, H-ZSM-23, KZ-1) or zeolites of the MFI structure type (for example, H-ZSM-5) or zeolites of the MEL structure type (for example, H-ZSM-11) or zeolites of the MTW structure type (for example, H-ZSM-12), or zeolites with the EUO structure type (for example, EU-1), or zeolite H-ZSM-57, or any member of the ferrierite structure family. Other examples of suitable catalysts are offretites, H-ZSM-4, H-ZSM-18 or zeolite Beta. Reference is made to "Synthesis of High-Silica Aluminosilicate Zeolites" by P. A. Jacobs and J. A. Martens (published as volume 33 in the series "Studies in Surface Science and Catalysis") for a review of the synthesis and properties of the aforementioned zeolites.

Additionally, the catalyst can be a zeolite synthesized without addition of a template, for example, faujasites, zeolite L, mordenites, erionites and chabazites, the structures of which are contained in the "Atlas of Zeolite Structure Types" by W. M. Meier and D. H. Olson (published by Butterworths on behalf of the Structure Commission of the International Zeolite Association). Zeolite catalysts having crystal structures that are essentially the same as the crystal structures of the above-mentioned zeolite catalysts but differ slightly therefrom in chemical composition may also be used, for example, zeolite catalysts obtained by removal of a number of aluminium ions from, or by steaming of, the above-mentioned zeolite catalysts, or zeolite catalysts obtained by addition of different elements, for example, by impregnation or cation exchange or by incorporation during the zeolite synthesis (for example boron, iron and gallium).

The zeolite catalyst may advantageously be a modified zeolite catalyst, as hereinafter defined, that has crystals of needle morphology, the ratio of the length to the diameter (L/D) of the crystals being not less than 3.

Preferably, H-ZSM-22, or a modified zeolite catalyst, as hereinafter defined, having the crystalline structure of ZSM-22 is used as the catalyst. Advantageously, there may be used crystals of H-ZSM-22 or of a modified zeolite catalyst, as hereinafter defined, having the crystalline structure of ZSM-22, the length of at least 75% of which do not exceed 10 μm and, more preferably not larger than 1.3 μ .

The process of the invention may be used with particular advantage in the oligomerisation of C₃-C₆-alkenes.

The water content of the alkene-containing feedstock is preferably from 0.08 to 0.25 molar %, more especially 0.1 to 0.2 molar %, based on the hydrocarbon content of the

feedstock. If, as may be desired, the alkene-containing feedstock contains as diluent a hydrocarbon gas other than a C₂-C₁₂-alkene, for example, a saturated hydrocarbon gas, that other hydrocarbon is to be included in the hydrocarbon content for the purposes of calculation of the water content.

Zeolite catalysts (other than modified zeolite catalysts, as hereinafter defined) may be made by any suitable method, for example, by heating a reaction mixture containing a source of silicon oxide, a source of aluminium oxide and, if appropriate, an organic promoter, for example, a nitrogen or phosphorus-containing organic base, together optionally with an alkali metal base, and separating the porous aluminosilicate crystals (zeolite precursor crystals) formed. The precursor crystals are then generally calcined in air or oxygen at a temperature exceeding 500°C, for example, at a temperature of 550°C for, for example, 10 to 20 hours. It is thought that, during calcination at such temperatures, the residual organic promoter remaining in the pores of the crystals is burnt substantially completely so that the pores are voided. The calcined material is preferably exchanged with ammonium ions (NH₄⁺) and subjected to conditions under which the ammonium ions decompose, with the formation of ammonia and a proton, thus producing the acidic form of the zeolite. Alternatively the acid form may be obtained by acid exchange with, for example, hydrochloric acid. If desired, however, the calcined material may be used as a catalyst without first being exchanged with ammonium ions, since the material then already possesses acidic sites. The activity of the material is then significantly lower than that of a material that has been exchanged with ammonium ions and then subjected to conditions under which the ammonium ions decompose.

The term "modified zeolite catalyst" is used throughout the specification, including the claims, to mean a zeolitic material that has been formed by a method in which an organic

substance (organic promoter or template) is used to promote formation of aluminosilicate crystals (zeolite precursor crystals) having the desired zeolitic structure, the uncalcined zeolite precursor crystals are exchanged with ammonium ions or protons, and the crystals are then calcined in conditions that are such that a portion of the organic promoter or of a decomposition product derived therefrom remains within the pores of the crystal.

Surprisingly, if needle-shaped aluminosilicate crystals (that is aluminosilicate crystals having needle morphology with a length to diameter (L/D) ratio of not less than 3) are exchanged with ammonium ions without first being subjected to the calcination step at a temperature exceeding 500°C customarily employed in the previously known methods, and then calcined under conditions such that the promoter remains, or decomposition products thereof remain, in the pores, the modified zeolite catalyst so obtained has a greater catalytic activity, giving good conversion with relatively little formation of cracked or saturated products or aromatics, and becomes deactivated in use more slowly, than material that has been calcined at temperatures exceeding 500°C prior to exchange with ammonium ions.

The present invention further provides a process for oligomerising C₂-C₁₂-alkenes comprising contacting a C₂-C₁₂-alkene-containing feedstock with a modified zeolite catalyst, as hereinbefore defined, having needle morphology with a L/D ratio of not less than 3 and a length not exceeding 30 µm. The modified zeolite catalyst is preferably one having the crystalline structure of ZSM-22.

The modified zeolite catalyst used in the processes described in the present specification may, for example, be prepared by

(i) heating a reaction mixture comprising

- (a) a source of silicon oxide (SiO_2)
(b) a source of aluminium oxide (Al_2O_3)
(c) optionally, an alkali metal source
(d) a nitrogen- or phosphorus-containing organic base,
5 and
(e) water;
- (ii) separating the formed crystals from the reaction mixture;
10
(iii) exchanging with ammonium ions or protons; and
(iv) calcining the crystals under conditions such that a portion of the organic base or of decomposition
15 products thereof remains within the crystal pores.

The organic base used in step (i) above may be, for example, a base containing a radical R_4N^+ in which the R groups may be the same or different and each may be hydrogen, an alkyl group
20 containing from two to eight carbon atoms or an aryl group containing up to eight carbon atoms. Preferably, however, the base is an amine, for example, an alkyl amine having from two to eight carbon atoms or an aryl amine having up to eight carbon atoms. More than one amine group may be present. For
25 example, for the manufacture of ZSM-22, the base may especially advantageously be 1,6-diaminohexane.

The exchange with ammonium ions in step (iii) may be carried out by any suitable method, for example, by treating the
30 crystals with an aqueous solution of ammonium chloride, ammonium nitrate or ammonium hydroxide. Hydrochloric acid treatment may be achieved by contacting the crystals with a dilute acid solution.

35 In step (iv), the crystals may be calcined at a temperature of from 120° to 430°C for from 5 hours to several days, generally at temperatures above 150°C . The temperatures in the upper part of the specified temperature range correspond to the

shorter heating times and the temperatures in the lower part of the specified temperature range correspond to the longer heating times.

5 Thus, for example, crystals may be calcined at a temperature of 400°C for from 5 to 20 hours. At a temperature of 120°C, longer calcination times of at least 2 days and preferably from 3 to 5 days will generally be necessary to achieve adequate voiding of the pores.

10

The calcination conditions should be chosen such that whereas sufficient of the organic base (template) is burnt that the pores are to a significant extent voided they still contain some base or decomposition products thereof. The presence of
15 the said base or decomposition products is indicated by a substantial discolouration of the crystals, as compared with the bright white material that is normally obtained after calcination at temperatures exceeding 500°C. Thus, for a given temperature, the calcination time should be so selected
20 that, while the crystals have a colour that is distinctly non-white, indicating the presence of decomposition products of the organic base, the pores are to a substantial extent voided. The extent to which the pores have been voided may be ascertained, for example, by measuring the net adsorption
25 capacity of the crystals using n-hexane. This may be done by contacting the crystals with nitrogen gas saturated with n-hexane at 30°C, and then monitoring the amount of n-hexane desorbed in a nitrogen atmosphere at the same temperature.

30

By way of example, the net adsorption capacity of the modified ZSM-22 used in accordance with the invention should preferably be reduced by not more than 50%, more preferably, by not more than 25%, as compared with the net adsorption capacity of ZSM-22 that has been obtained by a process in which calcination
35 has been carried out at 550°C for 16 hours both before and after exchange with ammonium ions. For other zeolite catalysts, the preferred maximum reduction in the net

adsorption capacity of the modified zeolite catalyst as compared with the corresponding unmodified catalyst may differ from that for ZSM-22, but the determination of preferred reductions for those other catalysts will be a matter for routine experiment.

The crystals may in some cases contain a minor proportion of crystalline material of a second aluminosilicate having a crystal structure that differs from the crystal structure of the aluminosilicate that forms the major proportion of the crystals, the proportion of the second aluminosilicate generally being sufficiently small that the properties of the crystal as a whole, including the external characteristics of the crystal, are determined predominantly by the aluminosilicate that constitutes the major proportion of the crystal. For example, a modified zeolite catalyst consisting of a major proportion of aluminosilicate material having the crystalline structure of ZSM-22, with co-crystallised aluminosilicate impurities having the crystalline structure of ZSM-5 and cristoballite, may be used.

The preferred molar ratios of the different reactants/silicon oxide source in the manufacture of H-ZSM-22 and the modified zeolite catalyst having the crystalline structure of ZSM-22 used in accordance with the present invention are as follows:

	Molar Ratio (MR)	MR (preferred)
SiO ₂ /Al ₂ O ₃	30 - 190	90
H ₂ O/SiO ₂	10 - 60	40
Template/SiO ₂	0.01 - 2.0	0.3
M ⁺ /SiO ₂	0 - 2.0	0.13 - 0.25

where M⁺ represents an alkali metal ion. The most preferred M⁺/SiO₂ ratio is 0.25.

Zeolite H-ZSM-22 used in accordance with the present invention preferably has a Si/Al molar ratio of from 20 to 75 and especially of approximately 35.

5 The zeolite catalysts or modified zeolite catalysts may be used in the form of powders (including powders consisting wholly or in part of single crystals). The zeolite catalysts may instead be incorporated in shaped agglomerates, for example, tablets, extrudates or spheres, which may be obtained
10 by combining the zeolite with a binder material that is substantially inert under the conditions employed in the oligomerisation process. The zeolite catalyst may be present in amounts of from 1 to 99% by weight, based on the combined weight of the zeolite and binder material. As binder material
15 there may be used any suitable material, for example, silica, metal oxides, or clays, such as montmorillonite, bentonite and kaolin clays, the clays optionally being calcined or modified chemically prior to use. Further examples of suitable matrix materials include silica-alumina, silica-beryllia, silica-
20 magnesia, silica-thoria, silica-titania, silica-alumina-magnesia, silica-alumina-thoria, silica-alumina-zirconia and silica-magnesia-zirconia.

In a further aspect of the invention there is provided a
25 process for oligomerising C₂-C₁₂-alkenes comprising contacting a C₂-C₁₂-alkene-containing feed-stock with crystals of H-ZSM-22 or with crystals of a modified zeolite catalyst having the crystalline structure of ZSM-22, the crystals having a length to diameter ratio of not less than 3 and a length of not
30 greater than 30 μm , preferably not greater than 10 μm and, more especially, not greater than 1 μm .

In an especially preferred process according to the invention, a C₂-C₁₂-alkene-containing feedstock, more especially a
35 feedstock containing C₃-C₆-alkenes, having a water content of from 0.05 to 0.25 molar % is contacted with H-ZSM-22 or a modified zeolite catalyst having essentially the crystalline

structure of ZSM-22. Especially good yields of dimeric, trimeric and tetrameric products are then obtained. Further, good catalyst activity is observed at relatively low temperatures, for example, at temperatures of from 150, typically 180, to 255°C. The proportions of alkanes and aromatics formed are low. The catalyst stability in the process is good in that the catalyst becomes deactivated relatively slowly, as compared with oligomerisation processes where the feedstock is relatively dry.

The zeolite catalysts used in accordance with the invention are regenerable. They are thermally and hydrothermally stable and, for example, can be regenerated by heating in air or oxygen at from 300 to 600°C or in steam at, for example, 500°C. The average number of branches per molecule of oligomer using H-ZSM-22 or a modified zeolite catalyst having the crystalline structure of ZSM-22 is relatively low compared with that obtained in previously known processes using solid phosphoric acid catalysts. For example nonenes produced from propylene using phosphoric acid catalysts have typically, about 2 branches, whereas those produced using zeolite catalysts have from 1.7 to 1.8 branches. Octenes produced using the phosphoric acid catalyst typically have 1.9 to 2 branches whereas with zeolite catalysis the branching can be lowered to 1.4. The lower branching leading to more reactive products.

The alkenes that may be oligomerised by the processes of the invention are ethene, propene, and linear or branched C₄-C₁₂-alkenes, which may be mono-, di- or polyunsaturated. As indicated above, however, the alkenes are preferably C₃-C₆-alkenes. The process is particularly advantageous for the oligomerisation of propene and butenes and may be used for the oligomerisation of a single alkene, or of mixtures of alkenes of the same or of different carbon numbers. The alkene may if desired be diluted with another suitable gas, for example, a low molecular weight saturated hydrocarbon.

The reaction is preferably carried out at pressures higher than atmospheric pressure, for example, at pressures up to 100 bar (10^7 Pa).

5 Certain illustrative embodiments of the invention will now be described with reference to the accompanying drawings, of which:

10 Fig. 1 is a graph showing % conversion of propene against catalyst life for H-ZSM-5;

Fig. 2a is a graph showing % conversion of propene against catalyst life for H-ZSM-22;

15 Fig. 2b is a graph showing % conversion of propene against catalyst life for a modified zeolite catalyst having the crystalline structure of ZSM-22;

20 Fig. 3 is a graph showing carbon number distribution of the product of butene oligomerisation; and

Fig. 4 is a graph showing the isomer distribution of C-8 product of butene oligomerisation.

25 In the following Examples, except where indicated to the contrary, NH_4Cl exchanged catalyst crystals had, prior to use, been calcined at a temperature of 550°C for from 15 to 20 hours in air both before and after exchange, oligomerisation was carried out without hydration of the feedstock, and
30 percentages are by weight except where indicated otherwise.

Example 1

Effect of hydration of feedstock on catalytic activity of H-ZSM-5, H-ZSM-12 and H-ZSM-22

5

(a) ZSM-22 precursor crystals having an average length of 1 μm were made by the method described in Example 4(a) below and the crystals were calcined in oxygen for 16 hours at 550°C, exchanged with NH_4Cl , and calcined again at 550°C for 16 hours in oxygen. A feedstock containing, on a hydrocarbon basis, 49.12% propene, 47.95% propane, the balance being ethane and butane was passed over H-ZSM-22 crystals. The product was analysed by gas chromatography (GC) at intervals. After 7 days, feedstock to be fed to the reactor containing the catalyst was hydrated by passing the feedstock, at a temperature of 39°C, through a thermostatted water saturator prior to its introduction into the reactor. The water content of the feedstock prior to its introduction into the water saturator was 0.02 molar % and, on leaving the saturator, the water content was 0.15 molar %, in each case, based on the hydrocarbon content of the feedstock. The pressure in the reactor was maintained at 70 bar (7×10^6 Pa) throughout. The results are summarised in Table 1A.

25

(b) A feedstock containing, based on hydrocarbon content, 49.42% propene, 50.50% propane, the balance being ethane and butane, and having an initial water content of 0.02 molar % based on the hydrocarbon content, was hydrated in the manner described in Example 1(a), to give a water content of 0.15 molar % and fed to a reactor containing H-ZSM-12 crystals. The pressure was maintained at 70 bar (7×10^6 Pa). The product was analysed. After 6 $\frac{1}{2}$ days the feed hydration apparatus was by-passed, feedstock being fed to the reactor without hydration, the unhydrated feedstock having a molar water

35

content of 0.02%, based on the hydrocarbon content. Further reaction conditions and the results are summarised in Table 1B below.

- 5 (c) A feedstock containing approximately 50% propene and 50% propane was hydrated in the manner described in Example 1(a) to give a water content of 0.15 molar % based on the hydrocarbon content and fed to a reactor containing H-ZSM-5 crystals. The reactor was maintained at a pressure of 70 bar (7×10^6 Pa) and, initially, at a temperature of 230°C. The temperature was subsequently raised to 250°C. The product was analysed at intervals. The experiment was then repeated using a feedstock having a water content of 0.02 molar % based on the hydrocarbon content of the feedstock. The results are shown in Fig. 1.

As is clear from Tables 1A and 1B and from Fig. 1 hydration of the feedstock provides significant advantages. Fig. 1 indicates that, for H-ZSM-5, the % conversion is increased significantly both at 230°C and 250°C. Table 1A shows that deactivation of H-ZSM-22 occurs relatively quickly when the feed is not hydrated. On commencement of hydration a significant improvement is observed, and an improved level of activity is maintained throughout the experiment subsequently. Thus, deactivation occurs less readily in cases where the feed has been hydrated than where no hydration is carried out. Further, the deactivation that has occurred during the use of unhydrated feed is even reversed by increasing the water content of the feed. A loss of activity is observed upon ceasing hydration of the feedstock in Example 1(b) - see Table 1B.

As will be seen from Tables 1A and 1B, the proportion of saturated reaction product is in each case not more than about 5%. In each case, the amount of aromatics formed was less than 50 ppm.

Table 1A

Hydration (°C)	NO HYD	NO HYD	NO HYD	NO HYD	NO HYD	NO HYD	NO HYD	39	39	39	39
Temperature (°C)	205	230	230	230	230	230	230	230	230	230	230
Space Vel. (weight/weight.hr)	1.26	1.62	1.44	1.40	1.60	1.59	1.28	1.50	1.19	1.35	1.35
Days on Stream	0.83	1.83	2.83	3.83	6.83	7.83	8.83	9.83	10.83	11.88	11.88
Catlife (weight product/ weight cat)	9.5	13.1	29.2	42.3	72.4	87.2	100.5	116.2	128.1	142.9	142.9
Propene Conversion (%)	76.78	18.69	94.91	79.97	53.11	78.99	88.08	88.87	84.41	89.83	89.83
Selectivity C5	0.56	0.29	0.64	0.48	0.38	0.51	0.51	0.52	0.30	0.59	0.59
C6	28.34	56.18	22.87	41.78	57.17	43.30	40.62	42.13	46.02	38.40	38.40
C7	1.35	1.06	2.32	1.50	0.98	1.32	1.31	1.34	1.41	1.56	1.56
C8	1.67	0.49	1.74	1.05	0.68	1.02	0.94	0.99	0.99	1.20	1.20
C9	35.06	26.78	32.33	31.37	28.05	31.79	32.02	31.34	31.69	31.85	31.85
C10	2.11	1.06	2.64	1.55	0.86	1.43	1.47	1.47	1.55	1.78	1.78
C11	1.91	0.67	2.39	1.27	0.60	1.18	1.20	1.15	1.30	1.49	1.49
C12	14.83	7.77	16.39	11.53	7.35	12.20	12.59	12.23	10.87	14.14	14.14
C13	2.76	0.00	3.62	1.94	0.00	0.97	1.60	1.39	0.00	2.16	2.16
C14	1.46	0.00	2.11	0.00	0.00	0.00	0.00	0.00	0.00	0.49	0.49
C15	9.94	5.70	12.94	7.51	3.92	6.28	7.74	7.43	5.87	6.35	6.35
C5-C15 Product from	11.82	3.57	15.47	7.81	3.51	6.43	7.04	6.86	5.54	9.27	9.27
Cracking (%) (*)											
Saturates in C5-C15 Product (%)	5.1	3.1	4.8	3.2	2.4	2.8	3.5	3.0			

(*) Sum of C5 + C7 + C8 + C10 + C11 + C13 + C14

Table 1B

Hydration (°C)	40	40	40	40	40	40	40	40	NO HYD	NO HYD	NO HYD
Temperature (°C)	180	180	150	170	170	170	180	180	180	180	180
Space Vel. (weight/Weight.hr)	1.90	1.83	1.99	1.56	1.81	1.97	2.14	1.95	1.97	1.97	1.97
Days on Stream	0.42	1.42	2.42	2.67	5.42	6.42	7.42	8.42	9.42	9.42	9.42
Catlife (weight product/weight cat)	9.3	30.3	44.5	48.6	96.7	117.4	139.1	153.7	162.1	162.1	162.1
Propene Conversion (%)	98.33	97.09	60.08	87.09	81.38	88.79	85.04	62.84	35.63	35.63	35.63
Selectivity C5	0.70	0.49	0.22	0.46	0.40	0.47	0.47	0.47	0.66	0.66	0.66
C6	2.61	2.38	4.75	2.88	3.27	3.17	3.98	8.30	17.35	17.35	17.35
C7	2.85	2.57	1.15	2.05	1.91	2.47	2.52	2.54	2.51	2.51	2.51
C8	4.60	4.36	2.24	3.87	3.66	4.40	4.34	3.84	3.07	3.07	3.07
C9	20.67	22.50	54.25	35.28	39.56	30.90	33.86	43.10	46.13	46.13	46.13
C10	5.00	4.74	2.69	4.22	3.95	4.49	4.35	3.50	2.84	2.84	2.84
C11	5.94	5.46	1.94	4.19	3.54	4.50	4.20	3.01	2.20	2.20	2.20
C12	26.39	26.88	20.80	26.18	24.72	25.91	24.80	20.20	14.96	14.96	14.96
C13	6.77	6.25	2.10	4.37	3.74	4.84	4.58	3.14	2.17	2.17	2.17
C14	6.00	5.66	1.62	3.55	3.04	4.13	3.86	2.50	1.64	1.64	1.64
C15	18.47	18.72	8.24	12.95	12.22	14.71	13.04	9.38	6.46	6.46	6.46
C5-C15 Product from Cracking (%) (*)	31.86	29.52	11.96	22.70	20.23	25.30	24.33	19.01	15.09	15.09	15.09
Saturates in C5-C15 Product (%)	4.7		2.2	3.8	2.9	3.5		3.3			

(*) Sum of C5 + C7 + C8 + C10 + C11 + C13 + C14

Example 2

Effects of hydration and temperature on activity of H-ZSM-22 and a modified zeolite catalyst having ZSM-22 crystalline structure.

- (a) Precursor crystals of ZSM-22 having an average size of 1 μm were prepared as described in Example 4(a) below and calcined at 550°C in oxygen for 16 hours, exchanged with aqueous NH_4Cl and calcined again at 550°C for 16 hours to obtain H-ZSM-22. 8.5 g of the H-ZSM-22 crystals so obtained were placed in a reactor. A feedstock consisting of, based on hydrocarbons, 50.68% propene, 48.52% propane, the balance being ethane and butanes, was hydrated by the method described in Example 1(a) to a moisture content of 0.15 molar %, based on the hydrocarbon content. The hydrated feedstock was passed over the catalyst at a temperature of 205°C and pressure of 70 bar (7×10^6 Pa). The space velocity was 1.50 to 2.10 weight/weight.hour. After the conversion had decreased to below 70%, the temperature was increased to 215°C, upon which the conversion increased to approximately 90%. On decrease of the conversion to below 80%, the temperature was increased to 225°C, the conversion then increasing to over 95%. At 225°C only a relatively slow decrease in conversion was observed. The effect of hydration was then investigated by ceasing hydration, the unhydrated feedstock having a water content of 0.02 molar % based on the hydrocarbon content. A rapid decrease in the conversion was observed. On recommencement of hydration the conversion immediately increased again. Analysis of the hexene isomer distribution showed the hexene product to contain, over the period of the reaction, from 11 to 13% linear hexenes, from 75 to 87% mono-branched hexenes and from 1 to 13% di-branched hexenes. Analysis of the nonene isomer distribution showed the nonene product to contain

1 to 4% linear nonenes, 14 to 40% mono-branched nonenes, 48 to 66% di-branched nonenes and 5 to 19% tri-branched nonenes. Fig. 2a shows the % conversion at a given catalyst life.

5

10

15

20

25

30

35

- (b) Precursor crystals of ZSM-22 having an average size of 1 μm were prepared as described in Example 4(a) below, exchanged with NH_4Cl and calcined in air at 400°C for 16 hours to obtain a modified zeolite catalyst having the crystalline structure of ZSM-22. 9.1 g of the calcined crystals were placed in a reactor. A feedstock containing, on a hydrocarbon basis, 50.70% propene, 48.32% propane, the balance being ethane and butanes, was hydrated by the method described in Example 1(a) to a moisture content of 0.15 molar % based on the hydrocarbon content. The hydrated feedstock was passed over the catalyst at a temperature of 205°C and with a pressure of 70 bar (7×10^6 Pa). The space velocity was from 1.60 to 2.20 weight/weight.hour. On observing a decrease in conversion towards 80%, the temperature was increased to 215°C , upon which the conversion increased to almost 100%. A relatively slow decrease in conversion was then observed and the conversion was once again increased to almost 100% by increasing the temperature to 225°C . The effect of hydration was examined by ceasing hydration, the feedstock being fed to the reactor having a water content of 0.02 molar % based on the hydrocarbon content. A rapid decrease in the conversion was then observed. Upon recommencement of hydration, however, the conversion was observed to increase dramatically. The results are shown in Fig. 2b. Analysis of the hexene isomer distribution in the product showed that the hexene fraction contained from 10 to 13% linear hexenes, 72 to 87% mono-branched hexenes and from 1 to 15% di-branched hexenes. The nonene fraction was found to contain from 2 to 4% linear nonenes, from 23 to 40% mono-branched

nonenes, from 50 to 62% di-branched nonenes and from 4 to 14% tri-branched nonenes.

Example 3

5

Oligomerisation of butene using a modified zeolite catalyst having the crystalline structure of ZSM-22

10

A modified zeolite catalyst having the crystalline structure of ZSM-22 was prepared as described in Example 2(b). A feedstock containing, based on the total hydrocarbon content, approximately 64% n-butenes and 35% butane, the balance consisting essentially of propane and propene, was hydrated as described in Example 1 to give a water content of

15

0.18 molar %, based on the hydrocarbon content. The hydrated feedstock was passed through 8.5 g of the modified zeolite catalyst in a reactor at a pressure of 70 bar (7×10^6 Pa) at a temperature which was gradually raised between 205 and 255°C. The space velocity was from 1.75 to 2.40

20

weight/weight.hour. The reaction product was analysed. The carbon number distribution of higher hydrocarbons, calculated for the cumulative product over a one month period, is shown in Fig. 3. Fig. 4 shows the isomer distribution for C₈ product (excluding saturated products). The conversion was above 80%.

25

Example 4

(a) Preparation of ZSM-22 crystals

30

144.03 g Ludox AS40 (which contains 40% by weight SiO₂ in water, with ammonium ions as stabilising agent) was dissolved in 248.01 g water. In a separate vessel, 6.90 g Al₂(SO₄)₃·18H₂O were mixed with 351.18 g water and stirred until a clear solution was obtained, to which were added 15.49 g potassium hydroxide (87.5% purity). The solution was stirred, 33.45 g 1,6-diaminohexane were

35

added, while continuing stirring, and the solution so obtained was added to the Ludox AS40 solution. Stirring was continued for five minutes. 787.1 g of the combined solution were placed in a stainless steel autoclave. The temperature was increased from room temperature to 160°C at a rate of 1°C per minute, and maintained at 160°C for 48 hours while stirring at 100 rpm., during which time crystals were formed. The product was recovered from the mother liquor, washed with water until the pH fell to 9.7 and dried overnight at 120°C. The ZSM-22 precursor crystals so obtained were shown by X-ray diffraction to consist of ZSM-22, containing traces of ZSM-5 impurities. The precursor crystals, which had a SiO₂/Al₂O₃ ratio of 63, were of needle morphology, having an average needle length of 1 µm and a L/D ratio of between 4 and 7.

Longer needles were obtained by, for example, increasing the temperature of crystallisation or increasing the water content of the synthesis mixture.

(b) Effect of H-ZSM-22 crystal size on catalyst activity

Precursor crystals having an average needle length of 27 µm, at least 75 out of every hundred of the crystals having a needle length of from 20 to 34 µm were prepared as described in Example 4(a) above, calcined at 550°C for 16 hours in oxygen, exchanged with NH₄Cl and calcined again at 550°C for 16 hours to obtain the acidic form of ZSM-22 (H-ZSM-22).

A feedstock consisting, on a hydrocarbon basis, of 50.68% propene and 48.52% propane, the balance being made up of ethane and butane, was hydrated, using the hydration method described in Example 1, the water content being increased from 0.02 to 0.15 molar %, based on the hydrocarbon content of the feedstock. The hydrated

feedstock was passed through the H-ZSM-22 crystals at a temperature of from 180°C, pressure of 70 bar (7×10^6 Pa) and space velocity of approximately 2 weight/weight.hour. The conversion of propene to higher molecular weight hydrocarbons was monitored by GC. When the conversion had decreased to 80%, the temperature was increased such that the conversion remained above 80%. In this way, the % conversion was maintained at 80% or more until a maximum temperature of 235°C was reached.

The catalyst life in weight product/weight catalyst was determined at the time at which the conversion became less than 80% at temperature 235°C. The catalyst life was determined in a similar manner for H-ZSM-22 crystals having average needle lengths of 27 μm , 15 μm , 7 μm and 1 μm , respectively. The catalyst lives of the crystals are given in Table 2.

In the case of the crystals having average length of 1 μm , the conversion remained at over 90% at temperature 225°C after two months use and the reaction was then terminated.

Table 2

Crystal Length μm^*	Catalyst Life (weight product/weight catalyst)
27 ± 7	252
15 ± 4	384
7 ± 2	636
1 ± 0.3	1240

* Based on SEM (scanning electron microscopy) analysis, at least 75 out of every hundred of the zeolite crystals were within the range given.

Example 5

Effect of temperature of calcination of zeolite catalyst precursor on catalytic activity of ZSM-22 and other, non-needle-shaped zeolites

(a) Precursor crystals of ZSM-12 were obtained by preparing a synthesis gel containing 0.41 M Na₂O, 0.89 M tetraethyl ammonium oxide, 0.04 M Al₂O₃, 10 M SiO₂ and 80 M water. After crystallisation in a static autoclave at 125°C for 12 days, the solid material formed was recovered, washed, and dried. The crystals were divided into two batches. The first batch was exchanged with aqueous NH₄Cl and heated in air to a temperature of 300°C for 16 hours, and the second batch was heated in air to 550°C, for 16 hours, exchanged with aqueous NH₄Cl, and calcined again for a further 16 hours at 550°C. 9.9 g of the first batch of crystals were then placed in a reactor under a pressure of 70 bar (7×10^6 Pa). A feedstock containing, based on the hydrocarbon content, 49.97% propene, 49.88% propane, the balance ethane and butane, and having a water content of 0.15 molar %, based on the hydrocarbon content of the feedstock, was passed over the catalyst at a space velocity of approximately 2 weight/ weight.hour. The % propene conversion was monitored by GC at 180°C and at 230°C. In each case less than 5 % conversion was observed. In contrast, using 9 g of the second batch of crystals with a feedstock containing, based on hydrocarbons, 49.42% propene, 50.50% propane, the balance ethane and butane, and having a moisture content of 0.15 molar % based on the hydrocarbon content, initial conversion of over 95% was observed at 180°C. In the case of the second batch, no measurements were made at 230°C.

(b) Precursor crystals of ZSM-22 and an average needle length of 27 μ m were prepared as described in Example 4(a) and

divided into two batches, the first batch being exchanged with aqueous HCl and calcined at 400°C to give a modified zeolite catalyst having the crystalline structure of ZSM-22, and the second batch being calcined at 550°C, exchanged with NH₄Cl, and calcined again 550°C.

Calcination was in each case carried out for 16 hours. 14.7 g of the first batch of crystals were placed in a reactor. A feedstock containing 49.20% propene, 47.95% propane, the balance ethane and butanes, based on hydrocarbon content, was fed to the reactor, after hydration in accordance with the hydration method described in Example 1 to give a water content of 0.15 molar %, based on the hydrocarbon content. The space velocity was from 1.45 to 2.05 weight/weight.hour.

The pressure in the reactor was maintained at 70 bar (7×10^6 Pa). The oligomerisation product was analysed. The procedure was then repeated using 15.4 g of the second batch of crystals, the feedstock containing, based on hydrocarbons, 48.70% propene, 48.42% propane, and having a water content of 0.15 molar % based on the hydrocarbon content of the feedstock, the balance ethane and butanes, and the space velocity being from 0.75 to 2.00 weight/weight.hour. The results are summarised in Table 3A for the first batch of crystals (modified zeolite catalyst) and in Table 3B for the second batch (H-ZSM-22). As shown in Tables 3A and 3B, the proportion of saturated C₅ to C₁₅ product is very small. The propene conversion decreases considerably more rapidly when H-ZSM-22 that has been calcined at 550°C is used than when the modified zeolite catalyst is used.

The amount of aromatics formed was in each case less than 50 ppm.

Table 3A

Space Vel. (weight/weight.hr)	1.94	2.01	1.51	1.50	1.46	1.54	1.56
Temperature (°C)	230	230	230	250	250	250	250
No Days on Stream	0.75	1.75	4.75	5.75	6.75	7.75	8.75
Catalyst Life (weight prod/weight cat)	17.1	40.9	91.1	106.7	121.2	135.0	148.5
Catalyst Life (weight feed/weight cat)	34.9	83.2	191.7	227.6	262.7	299.7	337.2
Propene Conversion (%)	99.78	99.90	94.10	88.36	83.66	76.28	72.72
Selectivity C5	1.09	0.94	0.47	0.34	0.27	0.22	0.20
C6	17.26	26.64	56.62	65.96	70.01	75.85	77.60
C7	3.04	2.30	1.05	0.83	0.66	0.57	0.54
C8	2.89	2.16	0.69	0.52	0.41	0.34	0.33
C9	20.73	26.67	26.26	22.84	20.72	17.53	16.49
C10	5.18	3.73	1.05	0.73	0.63	0.47	0.48
C11	5.15	3.46	0.98	0.65	0.51	0.41	0.38
C12	18.23	18.95	10.00	7.22	6.03	4.31	3.80
C13	5.64	3.39	0.69	0.00	0.00	0.00	0.00
C14	5.06	2.58	0.49	0.00	0.00	0.00	0.00
C15	15.71	9.18	1.71	0.91	0.75	0.30	0.18
C5-C15 Product from Cracking (%) (*)	28.06	18.56	5.42	3.07	2.49	2.01	1.92
Saturates in C5-C15 Product (%)	6.1	5.0					

(*) Sum of C5 + C7 + C8 + C10 + C11 + C13 + C14

Table 3B

Space Vel. (weight/weight.hr)	1.98	1.99	1.64	1.13	0.78	0.88	0.81
Temperature (°C)	230	230	230	250	250	250	250
No Days on Stream	0.75	1.75	4.75	5.75	6.75	7.75	8.75
Catalyst Life (weight prod/weight cat)	17.4	38.5	62.8	69.0	74.4	79.2	82.6
Catalyst Life (weight feed/weight cat)	35.8	83.7	201.9	229.1	248.3	267.9	287.4
Propene Conversion (%)	99.81	90.60	42.08	46.84	57.77	51.10	35.36
Selectivity (%)	1.14	0.97	0.29	0.26	0.25	0.30	0.22
C5	20.69	44.93	78.62	72.51	69.54	68.47	75.89
C6	3.21	1.80	0.69	0.83	0.75	0.76	0.64
C7	2.87	1.57	0.41	0.49	0.47	0.52	0.39
C8	24.53	29.57	15.46	19.65	21.83	21.84	17.99
C9	4.55	2.05	0.52	0.66	0.69	0.74	0.58
C10	4.35	2.09	0.43	0.58	0.56	0.63	0.45
C11	17.76	12.07	3.37	4.58	5.37	5.43	3.80
C12	4.49	1.49	0.00	0.00	0.00	0.45	0.00
C13	3.74	1.12	0.00	0.00	0.03	0.26	0.00
C14	12.66	2.34	0.19	0.43	0.49	0.61	0.04
C15							
C5-C15 Product from Cracking (%) (*)	24.36	11.09	2.35	2.82	2.77	3.65	2.29
Saturates in C5-C15 Product (%)	6.8	4.2					

(*) Sum of C5 + C7 + C8 + C10 + C11 + C13 + C14

Example 6**Comparison of activities and rates of deactivation of H-ZSM-22 and H-ZSM-5 in olefin oligomerisation**

(a) A feedstock consisting of, based on the hydrocarbon content, 48.48% propene, 48.37% propane, the balance consisting of ethane and butane, and having a water content of not more than 0.02 molar %, based on the hydrocarbon content, was passed through 9.7 g H-ZSM-5 crystals SiO₂/Al₂O₃ ratio 65) in a reactor at a flow rate of approximately 40 ml/hour. The temperature in the reactor was maintained at 230°C and the pressure at 70 bar (7×10^6 Pa). The content of the product was monitored using GC.

(b) The process of (a) was repeated except that the feedstock contained 49.12% propene and 47.95% propane and the zeolite H-ZSM-5 was replaced by 7.8 g H-ZSM-22 (SiO₂/Al₂O₃ ratio 65). In contrast to the H-ZSM-5 crystals used in (a), the H-ZSM-22 crystals are of needle-shaped configuration having a length to diameter ratio of greater than 3 and an average length of less than 30 μ .

As illustrated by the measurements given in Table 4, H-ZSM-5 is less active than H-ZSM-22 and becomes deactivated more quickly. Thus, at a catalyst life (weight feed/weight catalyst) of 93.4, the conversion of propene to higher molecular weight hydrocarbons had fallen to 17.75% for H-ZSM-5. For H-ZSM-22, at a catalyst life (weight feed/weight catalyst) of 98.6 the conversion is 94.91 and at a catalyst life of 247.5 the conversion had fallen only to 53.11%.

The amount of aromatics produced in the process of (b) was less than 50 ppm.

Table 4

Catalyst	ZSM-5	ZSM-5	ZSM-5	ZSM-22	ZSM-22	ZSM-22
Space Vel. (weight/weight.hr)	2.23	2.15	1.99	1.44	1.40	1.60
No Days on Stream	0.25	0.88	1.88	2.83	3.83	6.83
Catalyst Life (weight prod/weight cat)	4.3	8.1	12.2	29.2	42.3	72.4
Catalyst Life (weight feed/weight cat)	13.4	45.7	93.4	98.6	132.2	247.5
Propene Conversion (%)	65.42	24.45	17.75	94.91	79.97	53.11
Selectivity C ₅	0.65	0.71	0.71	0.64	0.48	0.38
C ₆	5.09	18.67	31.70	22.87	41.78	57.17
C ₇	2.70	2.17	1.75	2.32	1.50	0.98
C ₈	3.48	2.16	1.53	1.74	1.05	0.68
C ₉	37.88	45.52	42.23	32.33	31.37	28.05
C ₁₀	4.62	2.60	1.93	2.64	1.55	0.86
C ₁₁	4.71	2.35	1.65	2.39	1.27	0.60
C ₁₂	21.78	13.83	10.40	16.39	11.53	7.35
C ₁₃	5.08	2.77	1.99	3.66	1.94	0.00
C ₁₄	3.66	1.97	1.53	2.11	0.00	0.00
C ₁₅ ⁺	10.36	7.25	4.58	12.94	7.51	3.92
C ₅ -C ₁₅ Product from Cracking (%) (*)	24.89	14.73	11.09	15.47	7.81	3.51
Saturates in C ₅ -C ₁₅ Product (%)				4.8	3.2	2.4

(*) Sum of C₅ + C₇ + C₈ + C₁₀ + C₁₁ + C₁₃ + C₁₄

Claims:

1. A process for oligomerising C₂-C₁₂-alkenes to produce olefins of initial boiling point in the range 30 to 310°C, comprising contacting a C₂-C₁₂-alkene-containing feedstock having a water content of from 0.05 to 0.25 molar %, based on the hydrocarbon content of the feedstock, with a zeolite catalyst.
2. A process as claimed in claim 1, wherein the catalyst is selected from the following zeolites: zeolites of the TON, MTT, MFI, MEL, MTW, or EUO structure types, H-ZSM-57, zeolites of the ferrierite structure family, offretites, H-ZSM-4, H-ZSM-18, Zeolite Beta, faujasites, zeolite L, mordenites, erionites and chabazites.
3. A process as claimed in claim 1 or claim 2, wherein the catalyst is a zeolite catalyst, as hereinbefore defined, having crystals of needle morphology with a length:diameter ratio of not less than 3.
4. A process as claimed in claim 1 or claim 2, wherein the catalyst is H-ZSM-22 or a modified zeolite catalyst, as hereinbefore defined, having the crystalline structure of ZSM-22.
5. A process for oligomerising C₂-C₁₂-alkenes comprising contacting a C₂-C₁₂-alkene-containing feedstock with a modified zeolite catalyst, as hereinbefore defined, having crystals of needle morphology with length:diameter ratio of not less than 3.
6. A process as claimed in claim 5, wherein the modified zeolite catalyst has the crystalline structure of ZSM-22.
7. A process as claimed in any one of claims 1 to 6, wherein there is used a modified zeolite catalyst that has been

calcined at a temperature of from 120° to 430°C for a period of from 5 days to 5 hours.

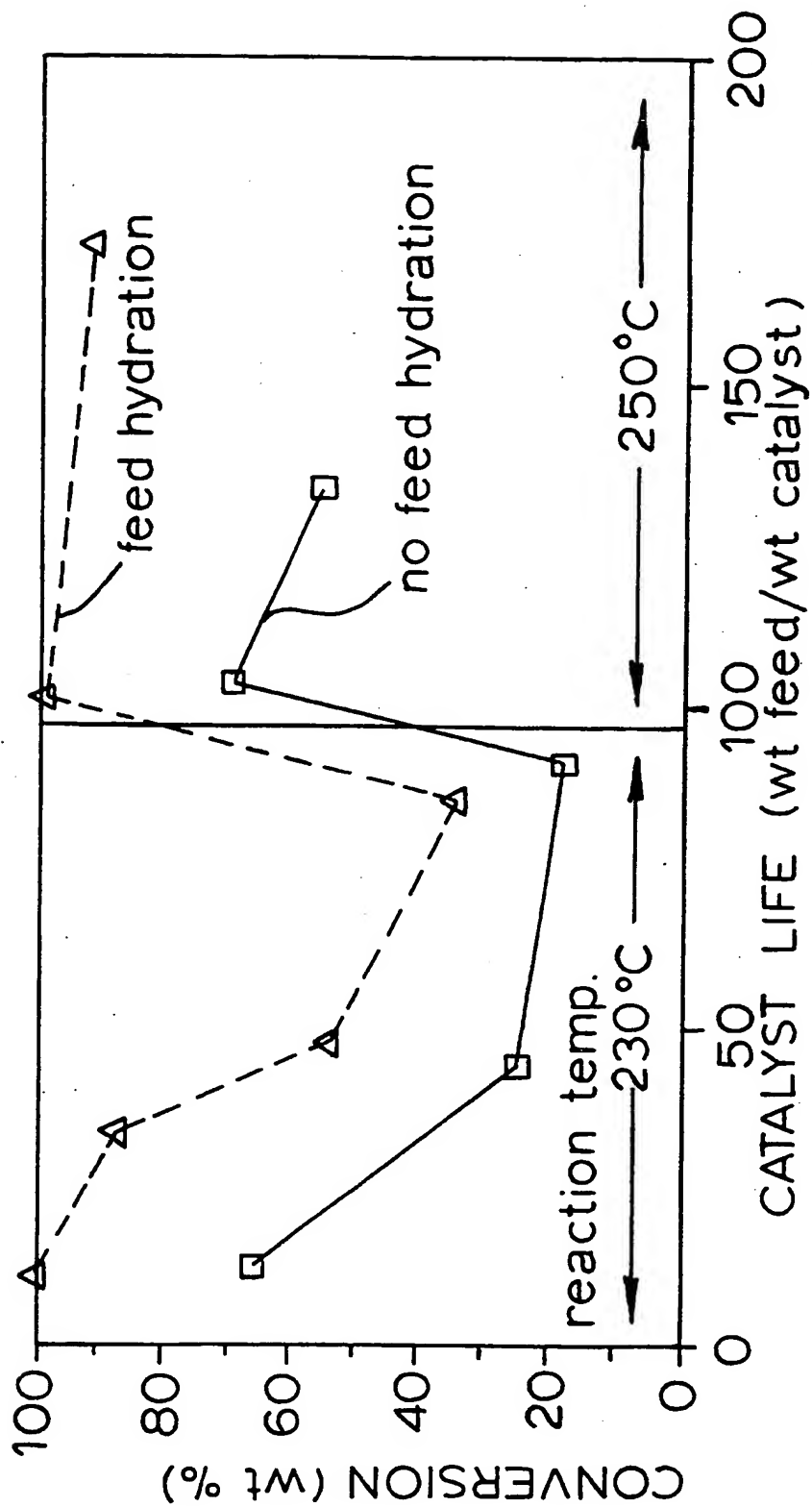
- 5 8. A process as claimed in any of claims 1 to 7, wherein the said contacting is carried out at a temperature not exceeding 260°C.
- 10 9. A process for oligomerising C₂-C₁₂-alkenes comprising contacting a C₂-C₁₂-alkene-containing feedstock with crystals of H-ZSM-22 or with crystals of a modified zeolite catalyst, as hereinbefore defined, having the crystalline structure of ZSM-22, at a temperature not exceeding 260°C.
- 15 10. A process as claimed in claim 9, in which the crystals have a length not exceeding 30 µm.
- 20 11. A process as claimed in claim 9 or claim 10, wherein the feedstock has a water content of from 0.05 to 0.25 molar %, based on the hydrocarbon content of the feedstock.
- 25 12. A process as claimed in any of claims 1 to 11, in which the feedstock contains C₃-C₆-alkenes, preferably propene or butene.
- 30 13. The use, for promoting catalyst activity and/or prolonging catalyst life in a zeolite-catalysed oligomerisation process, of a C₂-C₁₂-alkene-containing feedstock of from 0.05 to 0.25 molar % of water, based on the hydrocarbon content of the feedstock.
- 35 14. The use in the oligomerisation of C₂-C₁₂-alkenes of crystals having the porous aluminosilicate structure of ZSM-22 which have been formed using an organic promoter, for example, an alkyl amine or alkyl diamine, and which

contain residual promotor or decomposition product thereof.

- 5 15. The use, for increasing yields in the oligomerisation of C₂-C₁₂-alkenes, of a zeolite catalyst having needle morphology with a length:diameter ratio of not less than 3 and being in the form of crystals of length not exceeding 30 μm .

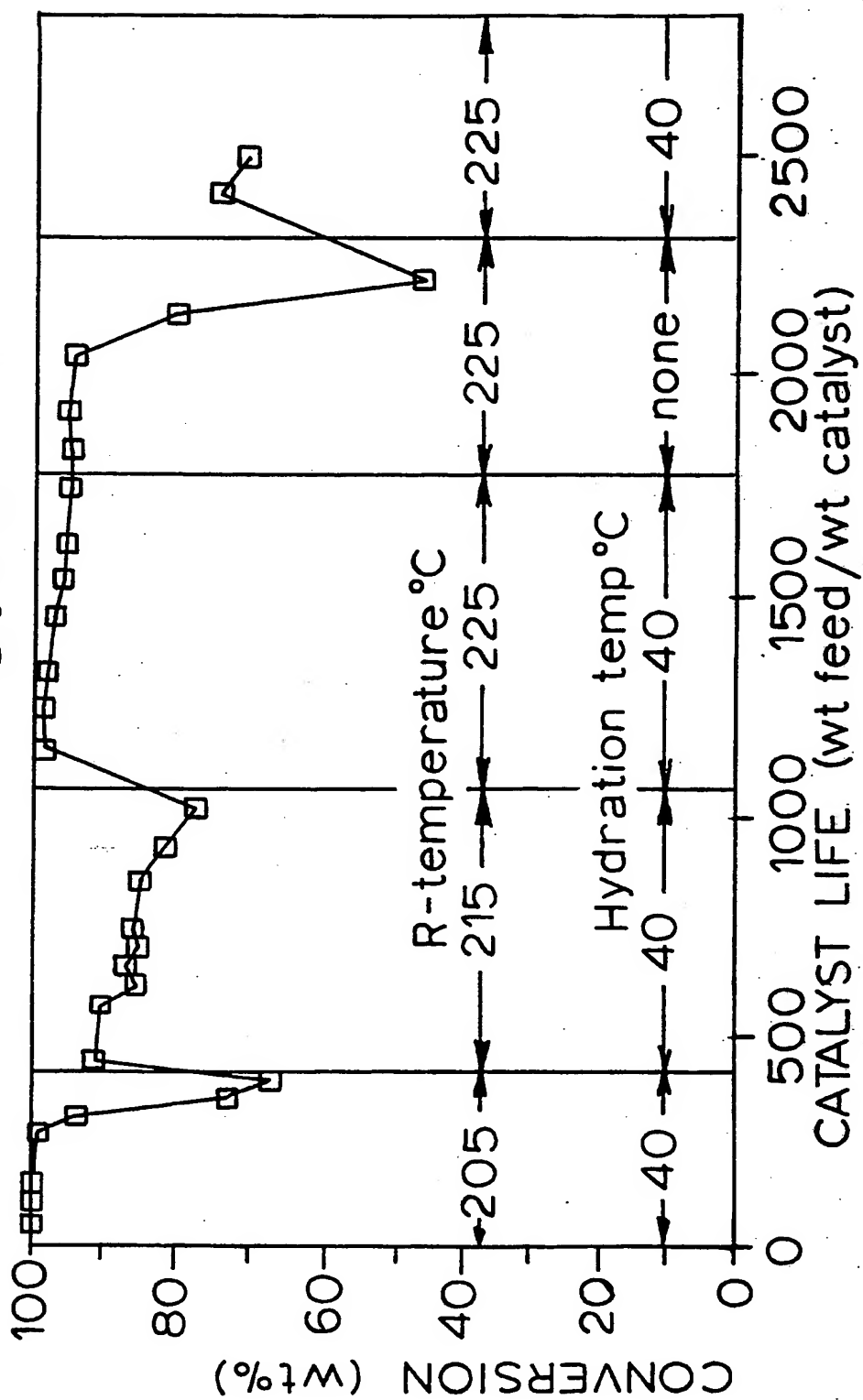
1/5

FIG.1



2/5

FIG. 2a



3/5

FIG. 2b

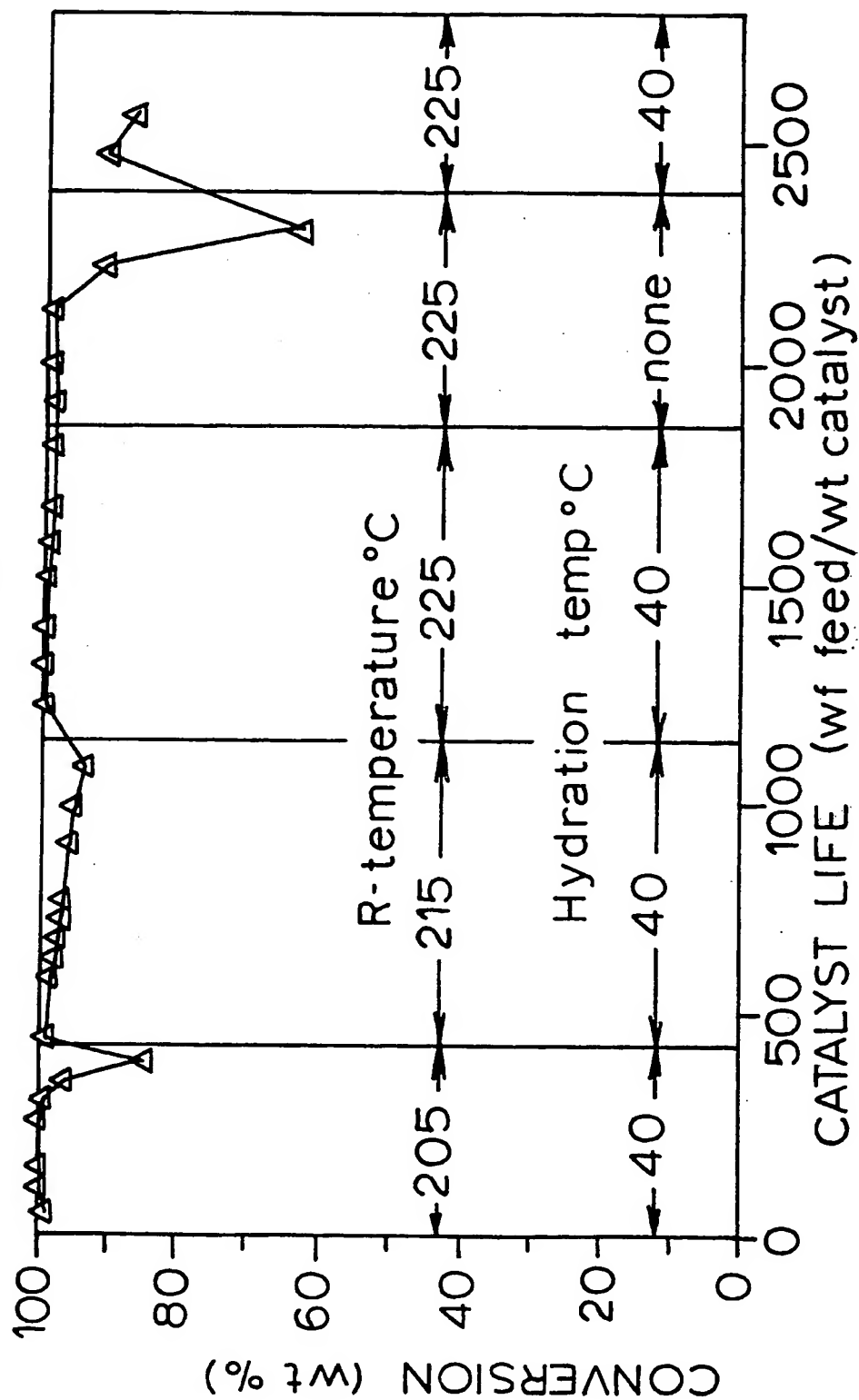
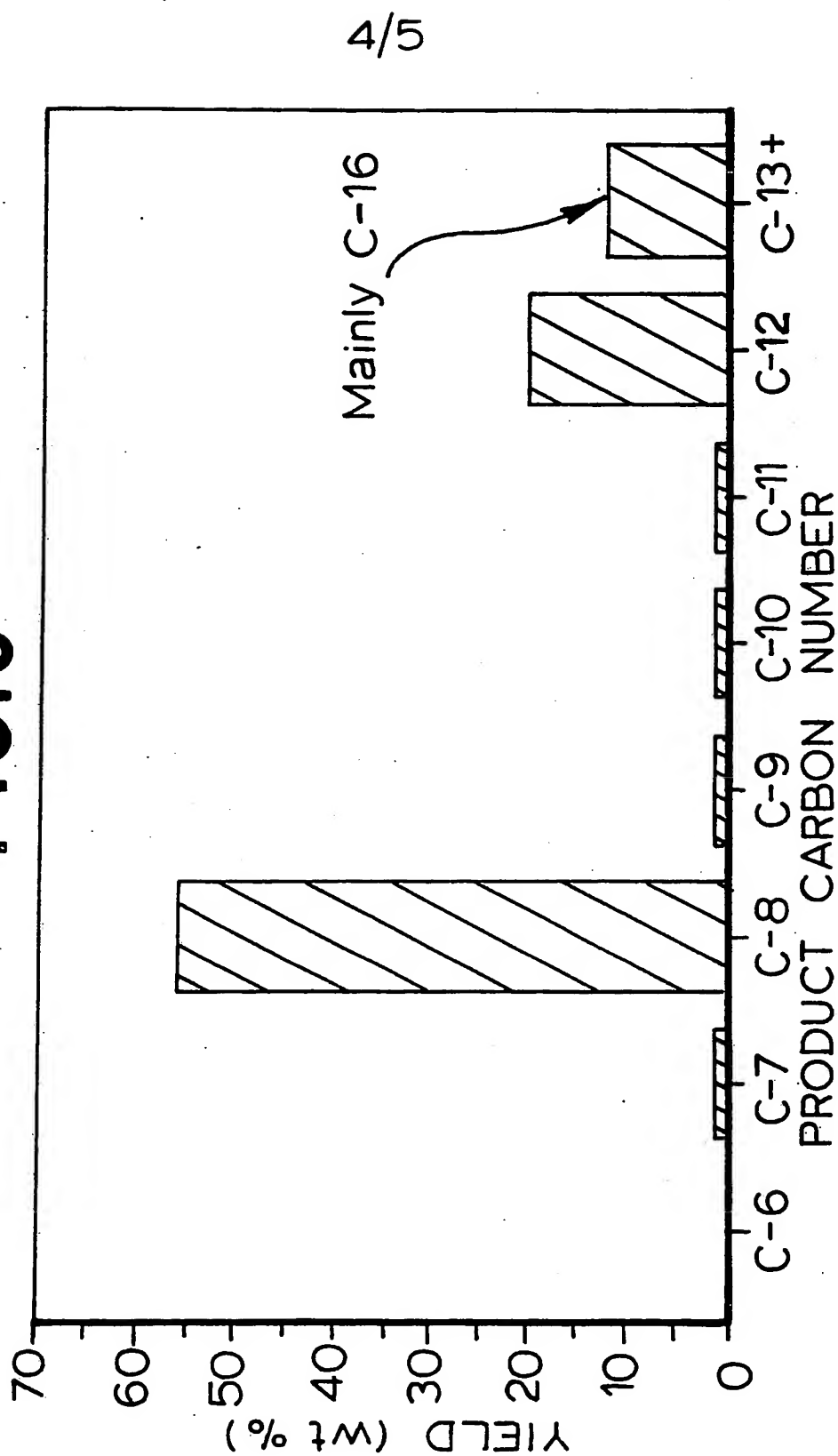


FIG. 3



5/5

